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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

First Named Inventor: Stenkamp, Dirk, et al) Art Unit: 1626

Application No.: 10/699,089) Examiner: Powers, Fiona

Filing Date: October 31, 2003) Confirmation No.: 1853

Title: Amide Compounds Having MCH-Antagonistic Activity and Medicaments Comprising these Compounds

Attorney Docket No.: 1/1407

IDS STATEMENT(S)



INFORMATION PREVIOUSLY CITED OR SUBMITTED

U.S. Application No. ("the parent application") is relied on by the above-identified application for an earlier effective filing date under 35 U.S.C. 120. The patents, publications, pending U.S. applications, or other information listed in the instant information disclosure statement were previously cited by the Office in the parent application or submitted to the Office in the parent application in an information disclosure statement that complied with 37 CFR 1.98(a)-(c). 37 CFR 1.98(d)



The fee set forth in 37 CFR 1.17(p) has been submitted herewith. 37 CFR 1.97(c-d) Otherwise, the Applicant authorizes the charging of all fees set forth in §1.16 and to §1.17 to Deposit Account No 02-29955 for the entire pendency of this application. 37 CFR 1.25(b)

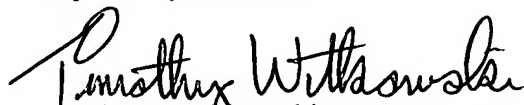


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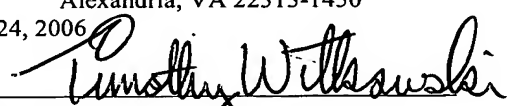
Each item of information contained in the instant information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three (3) months prior to the filing of the instant information disclosure statement. 37 CFR 1.97(e)(1)

- ☐ NO INFORMATION KNOWN MORE THAN THREE (3) MONTHS PRIOR
- No item of information contained in the instant information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing this certification after making reasonable inquiry, no item of information contained in the instant information disclosure statement was known to any individual designated in 37 C.F.R. §1.56(c) more than three (3) months prior to the filing of the instant information disclosure statement. 37 CFR 1.97(e)(2)
- ☐ NO INFORMATION RECEIVED FROM FOREIGN PATENT OFFICE MORE THAN THIRTY (30) DAYS PRIOR
- This submission contains only an information disclosure statement. Each item of information contained in the instant information disclosure statement was first cited in any communication from a foreign patent office in a counterpart application and this communication was not received by any individual designated in 37 C.F.R. §1.56(c) more than thirty (30) days prior to the filing of the instant information disclosure statement. 37 CFR 1.704(d)

Respectfully submitted,


Timothy X. Witkowski
Reg. No. 40,232

Patent Department
Boehringer Ingelheim Corp.
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Tel: (203) 798-4310

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PTO/SB/08a (08-03)

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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Sheet 1 of 10

Complete if Known

Application Number	10/699,089
Filing Date	October 31, 2003
First Named Inventor	Stenkamp, D., et al
Art Unit	1614
Examiner Name	To be assigned
Attorney Docket Number	1/1407

U. S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Document Number	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code ² (if known)			
		US- 3,994, 900	11/30/1976	Krapcho, John, et al	
		US- 6,366,268 B1	04/02/2002	Forrest, S.R., et al	
		US- 2002/0052383 A1	05/02/2002	Bakthavatchalam, R. et al	
		US- 2005/0026915 A1	02/03/2005	DeVita, R.J., et al	
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		US- 2004/0082780 A1	04/29/2004	Doherty, E.M., et al	
		US- 2003/0192132 A1	10/16/2003	Chassot, L., et al	
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FOREIGN PATENT DOCUMENTS

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
		WO 2005/063239 A1	07/14/2005	Boehringer Ingelheim International GmbH		
		WO 2005/085221 A1	09/15/2005	Boehringer Ingelheim International GmbH		
		DE 1 088 955	09/15/1960	Farbenfabriken Bayer AG		
		DE 3016827	11/20/1980	Continental Pharam S.A., Brussel		
		WO 2004/024702	03/25/2004	Boehringer Ingelheim Pharma GmbH		

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Sheet 2 of 10**Complete if Known**

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
		EP 1 593 667 A1	09/11/2005	Takeda Pharm. Co.		
		WO 2004/018467 A2	03/04/2004	BI Pharma GmbH		
		WO 2004/041820 A1	05/21/2004	BI Pharma GmbH		
		WO 03/045313 A2	06/05/2003	Merck & Co. Inc.		
		EP 1 283 199 A1	02/12/2003	Takeda Chem. Ind.		

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Sheet	3	of	10
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Application Number	10/699,089
Filing Date	October 31, 2003
First Named Inventor	Stenkamp, D., et al
Art Unit	1614
Examiner Name	To be assigned
Attorney Docket Number	1/1407

U. S. PATENT DOCUMENTS

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
		WO 01/55066 A2	08/02/2001	Syngenta Participat. In		
		WO 2004/039780 A1	05/13/2004	Boehringer Ingelheim		
		WO 02/04433 A2	01/17/2002	Neurogen Corporation		
		WO 03/013247 A1	02/20/2003	Syngenta Participat. In		
		WO 03/018579 A1	03/06/2003	Merck Frosst Canada		

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		WO 03/014111 A1	02/20/2003	Astrazeneca Ab		
		WO 02/28182 A1	04/11/2002	Syngenta Participation , Ag		
		WO 03/050087 A2	06/19/2003	Syngenta Participation Ag		
		WO 98/38156	09/03/1998	Takeda Chemical, Inc.		
		WO 03/024448 A2	03/27/2003	Methylgene, Inc.		

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Sheet 5 of 10

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
		EP 0 237 678 A1	09/23/1987	Glaxo Group Ltd.		
		WO 99/29674	06/17/1999	Janssen Pharmaceuticals		
		JP 04054118	02/21/1992	Tanabe Seiyaku Co.		
		WO 2004/072018	08/26/2004	Takeda Chemicals, Inc.		
		WO 2000/05223	02/02/2000	Zeneca Ltd., Inc.		

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		WO 03/033476 A1	04/24/2003	SmithKline Beecham Plc.		
		WO 99/02497 A2	01/21/1999	Novartis-Erfindungen		
		WO 02/092068	11/21/2002	Ono Pharmaceuticals		
		WO 03/045920 A1	06/05/2003	Merck & Co., Inc.		
		WO 03/049702	06/19/2003	Amgen, Inc.		

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Sheet	9	of	10	Attorney Docket Number	1/1407

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
		YANYUN CHEN, et al. "Targeted Disruption of the Melanin-Concentrating Hormone Receptor-1 Results in Hyperphagia and Resistance to Diet-Induced Obesity", Endocrinology 143(7):2469-2477 2002	
		DAQING QU, et al. "A role for melanin-concentrating hormone in the central regulation of feeding behaviour" Nature Vol. 380, pp 243-247 1996	
		MASAKO SHIMADA, et al. "Mice lacking melanin-concentrating hormone are hypophagic and lean" Nature Vol. 396, pp 670-674 1998	
		BETH BOROWSKY, et al. "Antidepressant, anxiolytic and anorectic effects of a melanin-concentrating hormone-1 receptor antagonist" Nature Medicine Vol. 8, No. 8, pp 825-830, 2002	
		DONALD J. MARSH, et al. "Melanin-concentrating hormone 1 receptor-deficient mice are lean, hyperactive, and hyperphagic and have altered metabolism" PNAS, Vol. 99, No. 5, pp 3240-3245, 2002	
		SHIRO TAKEKAWA, et al. "T-226296: a novel, orally active and selective melanin-concentrating hormone receptor antagonist" E. Journal of Pharm. Vol. 438, pp 129-13, 2002	
		J. KRAPCHO, et al; "Immunosuppressive Activity of 2'-(3-Dimethylaminopropylthio)cinnamanilide (Cinanserine) and Related Compounds" J. Med. Chemistry. 1969, 12(1), 164-166	
		TAMIZ, Amir P., et al; Structure-Activity Relationship of N-(Phenylalkyl)cinnamides as Novel NR2B Subtype-Selective NMDA Receptor Antagonists - J. Med. Chem. 1999, 42, 3412-3420	
		TETSUO, Sato, et al; Japanese Abstract 2002193800; VEGF Receptor antagonists for Treatment of neoangiogenesis-related diseases	

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¹ Applicant's unique citation designation number (optional). ² Applicant is to place a check mark here if English language Translation is attached.

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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Substitute for form 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(Use as many sheets as necessary)</i>				Complete if Known	
				<i>Application Number</i>	10/699,089
				<i>Filing Date</i>	October 31, 2003
				<i>First Named Inventor</i>	Stenkamp, D., et al
				<i>Art Unit</i>	1614
				<i>Examiner Name</i>	To be assigned
				<i>Attorney Docket Number</i>	1/1407
Sheet	10	of	10		

[illegible]

Examiner Signature		Date Considered	
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1 Applicant's unique citation designation number (optional). 2 Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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PTO/SB/17 (01-06)

Approved for use through 07/31/2006. OMB 0651-0032

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

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Fees pursuant to the Consolidated Appropriations Act, 2005 (H.R. 4818).

FEE TRANSMITTAL

For FY 2006

☐ Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT (\$)

Complete if Known

Application Number	10/699,089
Filing Date	October 31, 2003
First Named Inventor	Stenkamp, Dirk, et al
Examiner Name	Powers, Fiona
Art Unit	1626
Attorney Docket No.	1/1407

METHOD OF PAYMENT (check all that apply)☐ Check ☐ Credit Card ☐ Money Order ☐ None ☐ Other (please identify):☒ Deposit Account Deposit Account Number: 02-2955 Deposit Account Name: Boehringer Ingelheim Corporation

For the above-identified deposit account, the Director is hereby authorized to: (check all that apply)

☒ Charge fee(s) indicated below ☐ Charge fee(s) indicated below, except for the filing fee☒ Charge any additional fee(s) or underpayments of fee(s) under 37 CFR 1.16 and 1.17 ☒ Credit any overpayments

WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

FEE CALCULATION (All the fees below are due upon filing or may be subject to a surcharge.)**1. BASIC FILING, SEARCH, AND EXAMINATION FEES**

Application Type	FILING FEES		SEARCH FEES		EXAMINATION FEES		Fees Paid (\$)
	Fee (\$)	Small Entity Fee (\$)	Fee (\$)	Small Entity Fee (\$)	Fee (\$)	Small Entity Fee (\$)	
Utility	300	150	500	250	200	100	
Design	200	100	100	50	130	65	
Plant	200	100	300	150	160	80	
Reissue	300	150	500	250	600	300	
Provisional	200	100	0	0	0	0	

2. EXCESS CLAIM FEES

Fee Description	Fee (\$)	Small Entity Fee (\$)
Each claim over 20 (including Reissues)	50	25
Each independent claim over 3 (including Reissues)	200	100
Multiple dependent claims	360	180
Total Claims	Extra Claims	Fee (\$)
- 20 or HP =	x	=
HP = highest number of total claims paid for, if greater than 20.		
Indep. Claims	Extra Claims	Fee (\$)
- 3 or HP =	x	=
HP = highest number of independent claims paid for, if greater than 3.		

3. APPLICATION SIZE FEE

If the specification and drawings exceed 100 sheets of paper (excluding electronically filed sequence or computer listings under 37 CFR 1.52(e)), the application size fee due is \$250 (\$125 for small entity) for each additional 50 sheets or fraction thereof. See 35 U.S.C. 41(a)(1)(G) and 37 CFR 1.16(s).

Total Sheets	Extra Sheets	Number of each additional 50 or fraction thereof	Fee (\$)	Fee Paid (\$)
- 100 =	/ 50 =	(round up to a whole number) x	=	

4. OTHER FEE(S)

Non-English Specification, \$130 fee (no small entity discount)	Fees Paid (\$)
Other (e.g., late filing surcharge): Information Disclosure Statement	180.00

SUBMITTED BY

Signature		Registration No. 40,232	Telephone 203-798-4310
Name (Print/Type)	Timothy X. Witkowski		Date May 24, 2006

This collection of information is required by 37 CFR 1.136. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 30 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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